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AMENDMENTS TO THE CLAIMS

1-33. (Canceled).

34. (Previously presented) A method of enhancing a production of antibodies specific for a viral antigen, comprising:

identifying a subject in need of an enhanced production of antibodies specific for a viral antigen; and

providing to said subject an immunogenic composition comprising a viral antigen and ribavirin.

- 35. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is at least 0.25mg.
- 36. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is between about 0.25mg and 100 mg.
- 37. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is between about 0.25 mg and 25mg.
- 38. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is between about 0.25mg and 1mg.
- 39. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is at least 0.1 mg ribavirin per kg body weight of a subject receiving said composition.
- 40. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is at between about 0.1 mg ribavirin to about 1.0 mg ribavirin per kg body weight of a subject receiving said composition.
- 41. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is at between about 1.1 mg ribavirin to about 2.0 mg ribavirin per kg body weight of a subject receiving said composition.
- 42. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is at between about 2.1 mg ribavirin to about 3.0mg ribavirin per kg body weight of a subject receiving said composition.

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43. (Previously presented) The method of Claim 34, wherein the amount of ribavirin is at between about 3.1 mg ribavirin to about 4.0mg ribavirin per kg body weight of a subject receiving said composition.

- 44. (Currently amended) The method of Claim 34, wherein said <u>viral</u> antigen is obtained from a virus selected from the group consisting of hepatitis A virus, hepatitis B virus, and hepatitis C virus.
- 45. (Currently amended) The method of Claim 34, wherein said <u>viral</u> antigen is obtained from hepatitis C virus.
- 46. (Previously presented) A method of enhancing a production of antibodies specific for a viral antigen comprising:

providing an immunogenic composition comprising a viral antigen and ribavirin to a subject; and

measuring the production of antibodies specific for said viral antigen.

47. (Previously presented) The method of claim 46, wherein said measuring comprises measuring a reduction of viral load.

48-50. (Canceled)

51. (Previously presented) A method of increasing the titer of viral antigen-specific IgG antibodies in a subject in need thereof, comprising:

identifying a subject in need of an increase in titer of IgG antibodies that are specific for a viral antigen; and

providing said subject an immunogenic composition comprising ribavirin and said viral antigen.

- 52. (Previously presented) The method of claim 51, wherein said viral antigen is a hepatitis antigen.
- 53. (Previously presented) The method of claim 52, wherein said hepatitis antigen is an antigen from hepatitis A virus, hepatitis B virus, or hepatitis C virus.
- 54. (Previously presented) The method of claim 53, wherein said viral antigen is a hepatitis C virus antigen.
- 55. (Currently amended) The method of claim 54, wherein said viral antigen is an comprises a hepatitis C virus NS3 antigen.

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- 56. (Canceled)
- 57. (Previously presented) The method of claim 51, wherein the amount of ribavirin is between about 0.25mg and 100mg.
- 58. (Previously presented) The method of claim 51, wherein the amount of ribavirin is between about 0.25mg and 25mg.
- 59. (Previously presented) The method of claim 51, wherein the amount of ribavirin is between about 0.25mg and 1mg.
- 60. (Previously presented) The method of claim 51, wherein the amount of ribavirin is at least 0.1mg ribavirin per kg body weight of a subject receiving said composition.
- 61. (Previously presented) The method of claim 51, wherein the amount of ribavirin is between about 0.1mg ribavirin to about 1.0 mg ribavirin per kg body weight of a subject receiving said composition.
- 62. (Previously presented) The method of claim 51, wherein the amount of ribavirin is between about 1.1mg ribavirin to about 2.0 mg ribavirin per kg body weight of a subject receiving said composition.
- 63. (Previously presented) The method of claim 51, wherein the amount of ribavirin is between about 2.1mg ribavirin to about 3.0mg ribavirin per kg body weight of a subject receiving said composition.
- 64. (Previously presented) The method of claim 51, wherein the amount of ribavirin is between about 3.1mg ribavirin to about 4.0mg ribavirin per kg body weight of a subject receiving said composition.
- 65. (Previously presented) The method of claim 51, wherein the amount of ribavirin is at least 0.25mg.
- 66. (Previously presented) A method of enhancing a T cell response to a viral antigen in a subject in need thereof comprising:

identifying a subject in need of an improvement in a T cell response to a viral antigen; and

providing said subject an immunogenic composition comprising ribavirin and said viral antigen.

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- 67. (Previously presented). The method of claim 66, wherein said viral antigen is a hepatitis antigen.
- 68. (Previously presented) The method of claim 67, wherein said hepatitis antigen is an antigen from hepatitis A virus, hepatitis B virus, or hepatitis C virus.
- 69. (Previously presented) The method of claim 68, wherein said viral antigen is a hepatitis C virus antigen.
- 70. (Currently amended) The method of claim 69, wherein said viral antigen is an comprises a hepatitis C virus NS3 antigen.
 - 71. (Canceled)
- 72. (Previously presented) The method of claim 66, wherein the amount of ribavirin is at least 0.25mg.
- 73. (Previously presented) The method of claim 66, wherein the amount of ribavirin is between about 0.25mg and 100mg.
- 74. (Previously presented) The method of claim 66, wherein the amount of ribavirin is between about 0.25mg and 25mg.
- 75. (Previously presented) The method of claim 66, wherein the amount of ribavirin is between about 0.25mg and 1mg.
- 76. (Previously presented) The method of claim 66, wherein the amount of ribavirin is at least 0.1mg ribavirin per kg body weight of a subject receiving said composition.
- 77. (Previously presented) The method of claim 66, wherein the amount of ribavirin is between about 0.1mg ribavirin to about 1.0 mg ribavirin per kg body weight of a subject receiving said composition.
- 78. (Previously presented) The method of claim 66, wherein the amount of ribavirin is between about 1.1mg ribavirin to about 2.0 mg ribavirin per kg body weight of a subject receiving said composition.
- 79. (Previously presented) The method of claim 66, wherein the amount of ribavirin is between about 2.1mg ribavirin to about 3.0mg ribavirin per kg body weight of a subject receiving said composition.

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80. (Previously presented) The method of claim 66, wherein the amount of ribavirin is between about 3.1mg ribavirin to about 4.0mg ribavirin per kg body weight of a subject receiving said composition.

- 81. (New) The method of claim 51, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 200 consecutive amino acids.
- 82. (New) The method of claim 51, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 100 consecutive amino acids.
- 83. (New) The method of claim 51, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 50 consecutive amino acids.
- 84. (New) The method of claim 51, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 10 consecutive amino acids.
- 85. (New) The method of claim 66, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 200 consecutive amino acids.
- 86. (New) The method of claim 66, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 100 consecutive amino acids.
- 87. (New) The method of claim 66, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 50 consecutive amino acids.
- 88. (New) The method of claim 66, wherein said viral antigen is a fragment of hepatitis C virus NS3 protein that comprises at least 10 consecutive amino acids.